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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR		ATT	ORNEY DOCKET NO.
09/465,802	12/17/99	GARCIA-BLANCO		M 1	579-321
HM12/1106			コ	EXAMINER	
NIXON & VANDERHYDE PC ATTORNEY AT LAW				HUNT, J ART UNIT	PAPER NUMBER
1100 NORTH GLEBE ROAD STH FLOOR ARLINGTON VA 22201-4714				1642 DATE MAILED:	8

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

11/06/01



Office Action Summary

Application No. 09/465,802 Applicant(s)

Garcia-Blanco And Carstens

Examiner

Jennifer Hunt

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	The MAILING DATE of this communication appears	on the cover sheet with the correspondence address			
Period 1	for Reply				
	ORTENED STATUTORY PERIOD FOR REPLY IS SET MAILING DATE OF THIS COMMUNICATION.	TO EXPIRE3 MONTH(S) FROM			
af - If the	ter SIX (6) MONTHS from the mailing date of this communic	FR 1.136 (a). In no event, however, may a reply be timely filed ation. , a reply within the statutory minimum of thirty (30) days will			
co - Failui - Any i	mmunication. re to reply within the set or extended period for reply will, by	period will apply and will expire SIX (6) MONTHS from the mailing date of this statute, cause the application to become ABANDONED (35 U.S.C. § 133). mailing date of this communication, even if timely filed, may reduce any			
Status					
1) 💢	Responsive to communication(s) filed on Aug 28, 2	2001			
2a) 💢	This action is FINAL . 2b) ☐ This act	ion is non-final.			
3) 🗆	Since this application is in condition for allowance closed in accordance with the practice under <i>Ex pa</i>	except for formal matters, prosecution as to the merits is rte Quayle, 1935 C.D. 11; 453 O.G. 213.			
Disposi	tion of Claims				
4) 💢	Claim(s) <u>1-4</u>	is/are pending in the application.			
4	a) Of the above, claim(s)	is/are withdrawn from consideration.			
5) 🗆	Claim(s)	is/are allowed.			
. 6) 💢	Claim(s) <u>1-4</u>	is/are rejected.			
7) 🗌	Claim(s)	is/are objected to.			
8) 🗆	Claims	are subject to restriction and/or election requirement.			
Applica	tion Papers				
9) 🗆	The specification is objected to by the Examiner.				
10)	The drawing(s) filed on is/are	objected to by the Examiner.			
11)	The proposed drawing correction filed on	is: a) \square approved b) \square disapproved.			
12)	The oath or declaration is objected to by the Exami	ner.			
Priority	under 35 U.S.C. § 119				
13)□	Acknowledgement is made of a claim for foreign p	riority under 35 U.S.C. § 119(a)-(d).			
a) 🗀	☐ All b)☐ Some* c)☐ None of:				
	1. Certified copies of the priority documents hav	e been received.			
	2. \square Certified copies of the priority documents hav	-			
	3. Copies of the certified copies of the priority de application from the International Bure	au (PCT Rule 17.2(a)).			
14)	ee the attached detailed Office action for a list of th				
141	Acknowledgement is made of a claim for domestic	priority under 35 0.5.C. § 119(e).			
Attachm	ent(s)				
	otice of References Cited (PTO-892)	18) Interview Summary (PTO-413) Paper No(s).			
	otice of Draftsperson's Patent Drawing Review (PTO-948)	19) Notice of Informal Patent Application (PTO-152)			
17) Information Disclosure Statement(s) (PTO-1449) Paper No(s). 20) Other:					

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DETAILED ACTION

Continued Prosecution Application

1. The request filed on 9/28/2001 for a Continued Prosecution Application (CPA) under 37 CFR 1.53(d) based on parent Application No. 09/465,802 is acceptable and a CPA has been established. An action on the CPA follows.

Specification

2. It is noted that the specification contains numerous numerical citations, wherein a cited document is cited by a numeral, however there is no accompanying list which states what the reference numbers refer to. Thus numerous citations through the specification, particularly in the examples, fail to refer to the appropriate prior art documents.

Claim Rejections - 35 U.S.C. § 112

3. Claims 1-4 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claims 1-4 are unclear in the recitation of "metastatic potential", and "increase in metastatic potential". The metes and bounds of metastatic potential cannot be determined. It is not clear how such "potential" would be determined or measured. The qualities and properties

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encompassed by "metastatic potential" are never clearly set forth in the specification or claims. Further it is not possible how an increase in metastatic potential would be determined or measured. Specifically, it is not clear what level said increase is determined from (ie: increased above what?)

Claims 1-4 are unclear in the recitation of "predominant expression". The metes and bounds of "predominant expression" cannot be determined. It is not clear what would be considered predominant expression and what would not. Specifically, it is not clear at what point of difference would expression of one isoform over another be considered "predominant".

4. Claims 1-4 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for determination of prostate malignancy or androgen resistance in a rat patient or human cell line by assaying mRNA, does not reasonably provide enablement for determination of metastatic potential in patients including humans, or determination of metastatic potential by assaying the presence of the isoforms. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to practice the invention commensurate in scope with these claims.

Factors to be considered in determining scope and enablement are: 1) quantity of experimentation necessary, 2) the amount of direction or guidance presented in the specification, 3) the presence or absence of working examples, 4) the nature of the invention, 5) the state of the

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prior art, 6) the relative skill of those in the art, 7) the predictability of the unpredictability of the art, and 8) the breadth of the claims (see Ex parte Forman, 230 USPQ 546, BPAI, 1986).

The specification teaches measurement of FGF-R2 IIIb and IIIc isoform mRNA in 2 specific cell lines and subsequent determination, based solely on the properties of those cell lines, that the expression of FGF-R2 IIIc is correlative to increased likelihood of malignancy and androgen resistance. The specification also teaches production of antibodies to the FGF-R2 IIIb and IIIc isoform. The specification does not teach detection of the FGF-R2 IIIb and IIIc isoforms themselves, or detection of the FGF-R2 IIIb and IIIc isoforms using antibodies.

The claims are broadly drawn to determination of metastatic potential in any prostate cancer patient by measuring FGF-R2 IIIb and IIIc isoform mRNA or the isoforms themselves.

Those of skill in the art, recognize that expression of mRNA, specific for a tissue type, does not dictate nor predict the translation of such mRNA into a polypeptide. For example, Alberts et al. (Molecular Biology of the Cell, 3rd edition, 1994, page 465) teach that translation of ferritin mRNA into ferritin polypeptide is blocked during periods of iron starvation. Likewise, if excess iron is available, the transferrin receptor mRNA is degraded and no transferrin receptor polypeptide is translated. Many other proteins are regulated at the translational level rather than the transcriptional level. For instance, Shantz and Pegg (Int J of Biochem and Cell Biol., 1999, Vol. 31, pp. 107-122) teach that ornithine decarboxylase is highly regulated in the cell at the level of translation and that translation of ornithine decarboxylase mRNA is dependent on the secondary structure of the mRNA and the availability of eIF-4E, which mediates translation

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initiation. McClean and Hill (Eur J of Cancer, 1993, vol. 29A, pp. 2243-2248) teach that p-glycoprotein can be over expressed in CHO cells following exposure to radiation, without any concomitant over expression of the p-glycoprotein mRNA. In addition, Fu et al (EMBO Journal, 1996, Vol. 15, pp. 4392-4401) teach that levels of p53 protein expression do not correlate with levels of p53 mRNA levels in blast cells taken from patients with acute myelogenous leukemia, said patients being without mutations in the p53 gene. Thus, predictability of protein translation is not necessarily contingent on mRNA expression due to the multitude of homeostatic factors affecting transcription and translation. Thus one of skill in the art would not be able to determine metastatic potential of a prostate cell by measuring the isoforms using an antibody. Only mRNA methods are enabled. Any further extrapolation is an invitation to experiment, with correlation to diagnosis lacking as set forth above.

Further, it is also well established in the art that conclusions drawn in cell lines cannot easily be extended to human patients. Those of skill in the art recognize that in vitro assays are generally useful to screen the effects of agents on target cells. However, clinical correlations are generally lacking. The greatly increased complexity of the in vivo experiment as compared to the very narrowly defined and controlled conditions of an in vitro assay does not permit a single extrapolation of in vitro assays to human activity with any reasonable degree of predictability. Cancer is by it's very nature unpredictable, because it is the result of one or more of any number of alterations in normal cellular physiology. Further, in vitro cell lines are artificially generated

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models, conclusions from which cannot blindly be extended to human patients. Activity of a cancer cell line is not necessarily reflective of the actual in vivo activity of that type of cancer.

Therefor, for the reasons set forth above, one of skill in the art would not be enabled to practice the full scope of the invention as claimed.

Claim Rejections - 35 U.S.C. § 102

5. Claims 1-3 are rejected under 35 U.S.C. 102(a) as being anticipated by Carstens et al., Oncogene Vol. 15, pages 3059-3065, December 18, 1997.

Carstens et al. teaches a method of determining the increased metastatic potential (likelihood of malignancy and androgen resistance) of a prostate tumor in a patient (a mouse) by assaying for increased expression of the FGF-R2 IIIc isoform over the FGF-R2 IIIb isoform. The assay used measures mRNA transcripts. (See for example, abstract, and pages 3063-3064)

6. Claims 1-3 are rejected under 35 U.S.C. 102(b) as being anticipated by Yan et al., Molecular and Cellular Biology, Vol. 13, No. 8, pages 4513-4522, August, 1993.

Yan et al. teaches a method of determining the increased metastatic potential (likelihood of malignancy and androgen resistance) of a prostate tumor in a patient (a rat) by assaying for increased expression of the FGF-R2 IIIc isoform over the FGF-R2 IIIb isoform. The assay used measures mRNA transcripts. (See for example, abstract, and pages 4514 and 4517)

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Conclusion

7. This is a CPA of applicant's earlier Application No. 09/465,802. All claims are drawn to the same invention claimed in the earlier application and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the earlier application. Accordingly, THIS ACTION IS MADE FINAL even though it is a first action in this case. See MPEP § 706.07(b). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no, however, event will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

No claims are allowed.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jennifer Hunt, whose telephone number is (703) 308-7548. The examiner can normally be reached Monday through Thursday 6:30am to 5:00pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Anthony Caputa can be reached at (703) 308-3995. The fax number for the group is (703) 305-3014 or (703) 308-4242.

Communications via internet e-mail regarding this application, other than those under 35 U.S.C. 132 or which otherwise require a signature, may be used by the applicant and should be addressed to [anthony.caputa@uspto.gov].

All internet e-mail communications will be made of record in the application file. PTO employees do not engage in Internet communications where there exists the possibility that sensitive information could be identified or exchanged unless the record includes a properly signed express waiver of the confidentiality requirements of U.S.C. 122. This is more clearly set forth in the Interim Internet Usage Policy published in the Official Gazette of the Patent and Trademark on February 25, 1997 at 1195 OG 89.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the group receptionist, whose telephone number is (703) 308-0196.

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Jennifer Hunt

October 21, 2001